imarc

WE'LL EARN YOUR APPROVAL.

RESEARCH TRAINING & DEVELOPMENT

You Need to Reach the Finish Line. Like Anything Else, that Takes Training.

Your study may be short staffed. It may be operating on a tight budget. Yet one thing it cannot afford to be is derailed due to lack of experience or knowledge. Quality research training is in demand because the increasing needs of clinical research demand it. IMARC offers affordable training and development programs that will get your clinical research staff – coordinators, investigators, monitors, project managers and others – up to speed fast.

Training for Trials Worldwide

- Human Subject Protection
- GCP Compliance
- Introduction to Research Coordinating
- Monitoring 101
- GCP, JGCP and ICH Guidelines
- FDA Regulations:
  21 CFR 11, 50, 54, 56, 312 and 812

You no longer need to struggle with GCP compliance. An onsite IMARC Trainer can educate your team on applicable regulations. Before your next investigator's meeting, arrange for an IMARC Trainer to plan a General or Advanced GCP Refresher Course for meeting attendees. Need pocket training cards or customized tools developed? Done!

IMARC Trainers have prepared clinical research teams throughout the world for more than 10 years. Our approach to hands-on training empowers your team to competently handle any site requirement they put their hands on.

Be trained to:

- Implement the framework required to stay compliant.
- Navigate the regulatory maze with GCP training tailored specifically to your study team.
- Run your own investigator-initiated IDE or IND study with existing staff.
- Manage specific phases or processes while staying focused on achieving approval.

Shape Up Your Research Team

Expert Clinical Research Training for:

Monitors • Research Coordinators • Investigators • Project Managers

To learn more, contact John E. Lehmann, Director of Business Development **440.801.1540**

Stay Current in the Global Research Playing Field.

How do you ensure you are going to be – and remain – a relevant player? Start with a comprehensive orientation process from IMARC. Combine that with our high-quality continuing education programs. Finally, mix in the knowledge and experience our experts can share with your team to resolve issues and problem-solve any site management issue. Or, partner with IMARC and we will take care of it all for you as you train, develop and learn.

IMARC Trainers have contributed to drug and device research while ensuring data integrity and enforcing ethical standards for more than a decade. Our experience will be instrumental in helping you bring the latest advances in technology to those who need it as quickly as possible.

Training Sessions

Expand your regulatory knowledge of study operations with these courses:

- Introduction to Research Coordinating
- Monitoring "Musts"
- Understanding Clinical Research Sites *(for Project Managers and Monitors)*
- Clinical Site Management
- Human Subject Protection
- Monitoring Device Studies
- Device vs. Drugs
- Monitoring Team Roles for Global Studies
- Warning Letters… and the Implications for You

Training Techniques

IMARC training solutions are designed to provide you with a better understanding of your specific research setting and how it functions. These informative, interactive and inspirational solutions include:

- Didactic Presentations
- Group Activities
- Case Study Discussions
- Warning Letter Reviews & Discussions
- Games & Quizzes
- Virtual Study Experiences

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